

the basic classes of controlled substances listed.

Dated: September 14, 2009.

**Joseph T. Rannazzisi,**

*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Importer of Controlled Substances Notice of Registration

By Notice dated June 15, 2009, and published in the **Federal Register** on June 23, 2009 (74 FR 29718), Aptuit, 10245 Hickman Mills Drive, Kansas City, Missouri 64137, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Marihuana (7360), a basic class of controlled substance listed in schedule I.

The company plans to import a finished pharmaceutical product containing cannabis extracts in dosage form for packaging for a clinical trial study.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and § 952(a) and determined that the registration of Aptuit to import the basic class of controlled substance is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971, at this time. DEA has investigated Aptuit to ensure that the company's registration is consistent with the public interest. The investigation has included inspection and testing of the company's physical security systems, verification of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to 21 U.S.C. 952(a) and § 958(a), and in accordance with 21 CFR 1301.34, the above named company is granted registration as an importer of the basic class of controlled substance listed.

Dated: September 11, 2009.

**Joseph T. Rannazzisi,**

*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*

[FR Doc. E9-22451 Filed 9-17-09; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Importer of Controlled Substances Notice of Registration

By Notice dated June 15, 2009, and published in the **Federal Register** on June 24, 2009, (74 FR 30111), AllTech Associates Inc., 2051 Waukegan Road, Deerfield, Illinois 60015, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed in schedules I and II:

Drug	Schedule
Gamma Hydroxybutyric Acid (2010).	I
Heroin (9200) .....	I
Cocaine (9041) .....	II
Codeine (9050) .....	II
Hydrocodone (9193) .....	II
Meperidine (9230) .....	II
Methadone (9250) .....	II
Morphine (9300) .....	II

The company plans to import these controlled substances for the manufacture of reference standards.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and § 952(a) and determined that the registration of AllTech Associates, Inc. to import the basic classes of controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971, at this time. DEA has investigated AllTech Associates, Inc. to ensure that the company's registration is consistent with the public interest. The investigation has included inspection and testing of the company's physical security systems, verification of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to 21 U.S.C. 952(a) and § 958(a), and in accordance with 21 CFR 1301.34, the above named company is granted registration as an importer of the basic classes of controlled substances listed.

Dated: September 11, 2009.

**Joseph T. Rannazzisi,**

*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on July 2, 2009, GE Healthcare, 3350 North Ridge Avenue, Arlington Heights, Illinois 60004-1412, made application by renewal to the Drug Enforcement Administration (DEA) as a bulk manufacturer of Cocaine (9041), a basic class of controlled substance in schedule II.

The company plans to manufacture a radioactive product used in diagnostic imaging in the diagnosis of Parkinson's Disease and for manufacture in bulk for investigational new drug (IND) submission and clinical trials.

Any other such applicant, and any person who is presently registered with DEA to manufacture such a substance, may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrisette Drive, Springfield, VA 22152; and must be filed no later than November 17, 2009.

Dated: September 14, 2009.

**Joseph T. Rannazzisi,**

*Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.*

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Manufacturer of Controlled Substances Notice of Registration

By Notice dated June 3, 2009, and published in the **Federal Register** on June 9, 2009, (74 FR 27350), Norac Inc., 405 S. Motor Avenue, P.O. Box 577, Azusa, California 91702-3232, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Nabilone (7379), a basic class of controlled substance listed in schedule II.

The company plans to manufacture the bulk controlled substance for use in product development and for distribution to its customers.

No comments or objections have been received. DEA has considered the